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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 1 of 2

Complete if Known	
Application Number	10/806,899
Filing Date	3/23/2004
First Named Inventor	Petrou et al.
Art Unit	~~1614~~ 1634
Examiner Name	Stephen Thomas Kapushoc
Attorney Docket Number	1386/19

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Document Number Number - Kind Code[2] (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
/STK/	1	US-7,282,336	10-16-2007	Wallace et al.	
/STK/	2	US-7,078,515	07-18-2006	Wallace et al.	
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FOREIGN PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Foreign Patent Document Country Code[3] - Number[4] - Kind Code[5] (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T[6]
/STK/	3	WO 2005/014863	02-17-2005	Bionomics Limited		
/STK/	4	WO 2004/085674	10-07-2004	Bionomics Limited		
/STK/	5	WO 2002/050096	06-27-2002	Bionomics Limited		
/STK/	6	WO 2002/006521	01-24-2002	Bionomics Limited		

Examiner Signature	/Stephen Kapushoc/	Date Considered	08/04/2008

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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 2 of 2

	Complete if Known
Application Number	10/806,899
Filing Date	3/23/2004
First Named Inventor	Petrou et al.
Art Unit	~~1614~~ 1634
Examiner Name	Stephen Thomas Kapushoc
Attorney Docket Number	1386/19

NON PATENT LITERATURE DOCUMENTS

Examiner Initials*	Cite No.[1]	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T[2]
/STK/	7	Chou et al., "The lack of association between febrile convulsions and polymorphisms in SCN1A," Epilepsy Research, Vol. 54, pgs. 53-57 (2003).	
/STK/	8	Fujiwara et al., "Mutations of sodium channel a subunit type 1 (SCN1A) in intractable childhood epilepsies with frequent generalized tonic-clonic seizures," Brain, Vol. 126, pgs. 531-546 (2003).	
/STK/	9	Hirschhorn et al., "A comprehensive review of genetic association studies," Genetics in Medicine, Vol. 4, No. 2, pgs. 45-61 (2002).	
/STK/	10	Notification Concerning Transmittal of Copy of International Preliminary Report on Patentability for International Application No. PCT/AU2006/000841 dated January 3, 2008.	
/STK	11	Official Action for U.S. Patent Application Serial No. 10/482,834 dated August 2, 2007.	
/STK/	12	Official Action for U.S. Patent Application Serial No. 10/482,834 dated April 4, 2008.	
/STK	13	Official Action for U.S. Patent Application Serial No. 11/262,647 dated February 15, 2008.	
/STK	14	Ohmori et al., "Significant correlation of the SCN1A mutations and severe myoclonic epilepsy in infancy," Biochemical and Biophysical Research Communications, Vol. 295, pgs. 17-23 (2002).	
/STK/	15	Stafstrom et al., "Epilepsy Genes: The Link Between Molecular Dysfunction and Pathophysiology," Mental Retardation and Developmental Disabilitie Research Reviews, Vol. 6, pgs. 281-292 (2000).	
/STK/	16	Supplementary European Search Report corresponding to Australian Patent No. AU0200910 dated February 17, 2005.	

Examiner Signature	/Stephen Kapushoc/	Date Considered	08/04/2008

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